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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification ⁷ : A61K 31/4453, 31/12, 31/055, 31/00, A61P 37/00</p>		<p>A1</p>	<p>(11) International Publication Number: WO 00/69439 (43) International Publication Date: 23 November 2000 (23.11.00)</p>
<p>(21) International Application Number: PCT/DK00/00253</p>		<p>(81) Designated States: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p>	
<p>(22) International Filing Date: 12 May 2000 (12.05.00)</p>		<p>Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p>	
<p>(30) Priority Data: PA 1999 00659 12 May 1999 (12.05.99) DK</p>			
<p>(71) Applicant (for all designated States except US): NEUROSEARCH A/S [DK/DK]; 93 Pederstrupvej, DK-2750 Ballerup (DK).</p>			
<p>(72) Inventors; and</p>			
<p>(75) Inventors/Applicants (for US only): JENSEN, Bo, Skaaning [DK/DK]; Neurosearch A/S, 93 Pederstrupvej, DK-2750 Ballerup (DK). OLESEN, Søren, Peter [DK/DK]; Neurosearch A/S, 93 Pederstrupvej, DK-2750 Ballerup (DK). CHRISTOPHERSEN, Palle [DK/DK]; Neurosearch A/S, 93 Pederstrupvej, DK-2750 Ballerup (DK).</p>			
<p>(74) Common Representative: NEUROSEARCH A/S; Patent Department, 93 Pederstrupvej, DK-2750 Ballerup (DK).</p>			
<p>(54) Title: CHEMICAL COMPOUNDS HAVING ION CHANNEL BLOCKING ACTIVITY FOR THE TREATMENT OF IMMUNE DYSFUNCTION</p>			
<p>(57) Abstract</p>			
<p>The present invention relates to chemical compounds having inhibitory activity on an intermediate conductance Ca^{2+} activated potassium channel (IKCa), and the use of such compounds for the treatment or alleviation of diseases or conditions relating to immune dysfunction.</p>			

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CHEMICAL COMPOUNDS HAVING ION CHANNEL BLOCKING ACTIVITY FOR THE TREATMENT OF IMMUNE DYSFUNCTION

TECHNICAL FIELD

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The present invention relates to chemical compounds having inhibitory activity on an intermediate conductance Ca^{2+} activated potassium channel (IK_{Ca}), and the use of such compounds for the treatment or alleviation of diseases or conditions relating to immune dysfunction.

10

BACKGROUND ART

Ion channels are transmembrane proteins, which catalyse the transport of inorganic ions across cell membranes. The ion channels participate in processes as 15 diverse as the generation and timing of action potentials, synaptic transmissions, secretion of hormones, contraction of muscles, etc.

Many drugs exert their effects via modulation of ion channels. Examples are anti-epileptic compounds like Phenytoin and Lamotrigine, which block voltage dependent Na^+ -channels in the brain, anti-hypertensive drugs like Nifedipine and 20 Diltiazem, which block voltage dependent Ca^{2+} -channels in smooth muscle cells, and stimulators of insulin release like Glibenclamide and Tolbutamide, which block an ATP-regulated K^+ -channel in the pancreas.

There is a large and still growing demand for non-toxic immune-regulating agents for use in relation to e.g. organ transplantation and auto-immune diseases.

25 Some of the currently used immune-suppressive compounds such as Cyclosporin A and FK506 prevent immunological proliferation by inhibition of the Ca^{2+} /calmodulin-dependent Ser/Thr phosphatase calcineurin. The usefulness of this class of compounds is limited by their side effects such as renal dysfunction, arterial hypertension, neurological effects (headache, insomnia, tremors, parasthesias, 30 lethargy), gastrointestinal effects (nausea, vomiting, diarrhoea), and diabetes.

Another class of compounds comprising e.g. Azathioprine and Mizoribine interfere in a cytotoxic manner directly with the DNA-replication process. Although cytotoxicity shows some selectivity towards strongly proliferating cells such as activated T- and B-lymphocytes, complications may follow due to effects on dividing 35 cells in the entire body, including bone marrow, hair sacs, the skin, testis, ovary and epithelia such as the airways, the intestinal tract, and the thick ascending limb of the loop of Henle's.

A fairly new approach for suppression of immune responses is to interfere with ion channels in the plasma membrane of cells in the immune system, especially

the T- and B-lymphocytes. Upon exposure to antigens by antigen presenting macrophages or to mitogens such as IL-2 or IFN- γ , an initial signal in the switching from the resting phase to the proliferating phase is an activation of the phosphoinositide signalling pathway resulting in an increase in the intracellular 5 concentration of Ca^{2+} ($[\text{Ca}^{2+}]_i$) due to Ca^{2+} release from intracellular stores. A sustained elevated $[\text{Ca}^{2+}]_i$ is maintained by an increased passive influx through mitogen regulated, voltage-independent Ca -channels. This increase in $[\text{Ca}^{2+}]_i$ is vital for the subsequent events leading to cell proliferation and secretion of lymphokines.

In resting T- and B-lymphocytes, the $[\text{Ca}^{2+}]$ is approximately 10^7 fold higher 10 outside versus inside the cell, and the membrane potential is negative inside, i.e. there is an inwardly directed electrochemical Ca^{2+} gradient. Thus, when the Ca -channels are activated they conduct Ca^{2+} into the cell. However, Ca^{2+} influx via the Ca -channels, tends to reduce or even eliminate this gradient, and thus to reduce the influx. Concomitant opening of K-channels keeps the membrane potential negative, 15 and activation of these channels is therefore essential for maintaining a large inwardly directed, electrochemical driving force for Ca^{2+} .

In the presence of blockers of lymphocyte K-channels, the cells depolarise, and thereby the Ca^{2+} influx necessary for the activation of the immune response is reduced.

20 Several types of K-channels have been described in B- and T-lymphocytes including both voltage-dependent K-channels (K_v), and voltage-independent Ca^{2+} -activated K-channels (K_{Ca}). It is well established, that the K_v -channels are activated by the Ca^{2+} -induced depolarisation of the lymphocyte, and non-selective blockers of K_v -channels are therefore quite effective immune-suppressive agents. However, these 25 compounds in general have severe side effects due to block of repolarization in excitable tissue (seizures, myotonic runs, high blood pressure, etc.).

Considerable effort has been made into the development of immune-selective K_v -blockers. The molecular rationale for this, has been the observation that T-lymphocytes express homomeric $K_v1.3$ -channels in contrast to excitable cells, 30 which always express several heteromeric subtypes of the K_v -channels.

A selective blocker of the $K_v1.3$ -homomer might therefore be an ideal, relatively non-toxic, immune-suppressive agent. Initial reports from these pharmacological programs indicate that selective $K_v1.3$ -blockers are very effective as anti-inflammatory agents. However, the well-known toxicity of non-selective K_v -blockers has apparently not disappeared. An example is the potent $K_v1.3$ blocker CP-35 35 339,818. This compound is also a potent blocker of $K_v1.4$, a cardiac and neuronal A-type K-channel. The side-effect of this compound is predicted to be interference with

the cardiac action potential (long QT-syndrome toxicity) as well as with the action potential repolarization and after hyperpolarization in neurons.

WO 97/34589 describes triaryl methane compounds that inhibit mammalian cell proliferation, inhibit the Gardos channel of erythrocytes, reduce sickle erythrocyte 5 dehydration and/or delay the occurrence of erythrocyte sickling or deformation, and suggest the use of these compounds in abnormal cell proliferation. However, the effect of these compounds on human T cell proliferation, the use of such compounds in normal cell proliferation as immune-suppressive agents, as well as their unexpected properties when used in combination therapy has never been disclosed.

10

SUMMARY OF THE INVENTION

A hitherto untested alternative to the block of the voltage-dependent K-channels is a selective inhibition of the Ca^{2+} -activated K-channels in T- and B- 15 lymphocytes. These channels are directly activated by the increased $[\text{Ca}^{2+}]_i$ which is the primary signal for lymphocyte activation. Further, contrary to K_v -channels, these channels are voltage-independent, and therefore they do not close upon hyperpolarization, implicating that they are even more effective than K_v channels in maintaining a large inward driving force on Ca^{2+} under conditions of elevated 20 intercellular Ca^{2+} -concentrations.

Two types of Ca^{2+} -activated K-channels have been described from lymphocytes: 1) Small-conductance, apamin-sensitive, Ca^{2+} -activated K-channels (SK_{Ca}) and 2) Intermediate-conductance, inwardly rectifying, Clotrimazole-sensitive, Ca^{2+} -activated K-channels (IK_{Ca}), also referred to as Gardos-channels. Resting T- 25 lymphocytes express both SK_{Ca} and IK_{Ca} , whereas B-lymphocytes only express IK_{Ca} .

Upon activation, prior to cell proliferation, the expression level of IK_{Ca} increases approximately 30 fold in both T- and B-lymphocytes. The expression levels of both $K_v1.3$ and SK_{Ca} remain unchanged, indicating a major role for the IK_{Ca} -channel in induction of T- and B-cell proliferation. Contrary to the SK_{Ca} -channels, which are 30 extensively expressed in CNS and heart (measured as mRNA abundance by Northern hybridisation) and in PNS, skeletal muscle, hepatocytes (measured as functional channels by electrophysiology), expression of IK_{Ca} -channels have never been reported from any excitable tissue. In fact, blood cells such as erythrocytes, monocytes, lymphocytes, endothelial cells, and certain cell-lines with an epithelial 35 ancestry, Ehrlich ascites tumour cells and HeLa cells appear to be the main source of this type of channels.

Furthermore, the very recent cloning of IK_{Ca} has enabled the demonstration of the mRNA for this gene in several organs including placenta, salivary glands, lung and pancreas. Thus, specific blockers of IK_{Ca} are likely to be very effective as

immune-suppressive agents, and devoid of side effects on excitable tissue. In fact, the IK_{Ca} -inhibitor Clotrimazole (which is also a blocker of the cytochrome P-450 system) has been extensively used clinically in the systemic treatment of fungal infections. No toxicity related to K-channel blockade has been described.

5 Accordingly, in its first aspect, the invention relates to the use of a chemical compound having IK_{Ca} inhibitory activity for the manufacture of a medicament for the treatment or alleviation of diseases, disorders or conditions relating to immune dysfunction.

10 In another aspect the invention provides a pharmaceutical compositions for use in the treatment or alleviation of diseases, disorders or conditions relating to immune dysfunction, comprising an effective amount of a chemical compound having IK_{Ca} inhibitory activity.

DETAILED DISCLOSURE OF THE INVENTION

15

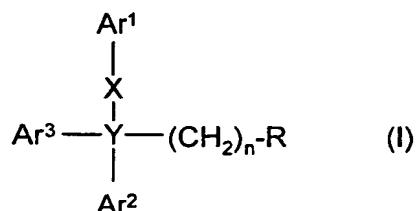
The present invention relates to the use of a chemical compound having selective IK_{Ca} inhibitory activity for treatment or alleviation of diseases or conditions relating to immune dysfunction.

20 Chemical Compound having IK_{Ca} Inhibitory Activity

According to the invention, chemical compounds having selective IK_{Ca} inhibitory activity may be identified by its ability to inhibit current through an IK_{Ca} channel, while showing essentially no effect at other potassium channels at a 10 fold higher concentration, as determined by conventional patch clamp technique.

25 The compounds for use according to the invention show IK_{Ca} inhibitory activity in concentrations below 100 μ M, preferably below 10 μ M, more preferred below 1 μ M. In its most preferred embodiment compounds show IK_{Ca} inhibitory activity show activity in low micromolar and the nanomolar range.

30 In a preferred embodiment the chemical compounds for use according to the invention showing selective IK_{Ca} inhibitory activity are triaryl methane derivatives represented by the general Formula I



and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5 or 6;

X is absent, or represent a group of the formula $-(CH_2)_n-$, of the formula

5 $-(CH_2)_n-Z-$ (in either direction), of the formula $-(CH_2)_n-CH=N-$ (in either direction), the formula $-(CH_2)_n-Z-(CH_2)_m-$, or of the formula $-(CH_2)_n-CH=N-(CH_2)_m-$ (in either direction), or a group of the formula $-R'''C(O)N-$;

in which formulas n and m, independently of each another, represent 0, 1, 2, 3 or 4; and Z represents O, S, or NR''' , wherein R''' represents hydrogen or alkyl;

10 Y represents a carbon atom (C), a nitrogen atom (N), or a phosphor atom (P), a silicium atom (Si), or a germanium atom (Ge);

Ar¹, Ar² and Ar³, independently of each another, represents a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more 15 times with substituents selected from the group consisting of halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula $-OR'$, $-SR'$, $-R'OR'$, $-R'SR'$, $-C(O)R'$, $-C(S)R'$, $-C(O)OR'$, $-C(S)OR'$, $-C(O)SR'$, $-C(S)SR'$, $-C(O)NR'(OR')$, $-C(S)NR'(OR')$, $-C(O)NR'(SR')$, $-C(S)NR'(SR')$, $-CH(CN)_2$, $-C(O)NR'_2$, $-C(S)NR'_2$, $-CH[C(O)R']_2$, $-CH[C(S)R']_2$, $-CH[C(O)OR']_2$, $20 -CH[C(S)OR']_2$, $-CH[C(O)SR']_2$, $-CH[C(S)SR']_2$, $-CH_2OR'$, or $-CH_2SR'$;

R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula $-OR'$, $-SR'$, $-R'OR'$, $-R''SR'$, $-C(O)R'$, $-C(S)R'$, $-C(O)OR'$, $-C(S)OR'$, $-C(O)SR'$, $-C(S)SR'$, $-C(O)NR''(OR')$, $-C(S)NR''(OR')$, $-C(O)NR''(SR')$, $-C(S)NR''(SR')$, $-CH(CN)_2$, $-C(O)NR'_2$, $-C(S)NR'_2$, $25 -CH[C(O)R']_2$, $-CH[C(S)R']_2$, $-CH[C(O)OR']_2$, $-CH[C(S)OR']_2$, $-CH[C(O)SR']_2$, $-CH[C(S)SR']_2$, $-CH_2OR'$, or $-CH_2SR'$; or a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected 30 from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula $-OR'$, or $-SR'$; and

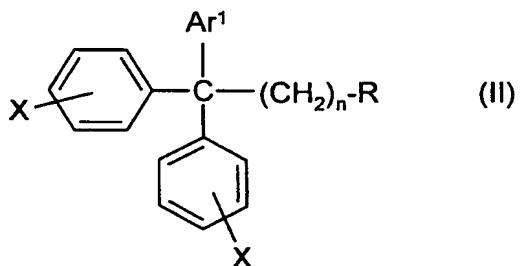
R' and R'', independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

In a more preferred embodiment, the triaryl methane derivative for use according to the invention is represented by the general Formula I wherein;

35 the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, 5 pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

In another preferred embodiment the triaryl methane derivative for use according to the invention is represented by the general Formula II



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and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5 or 6;

Ar¹ represents a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may 15 optionally be substituted one or more times with substituents selected from the group consisting of halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'', -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), 20 -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)₂, -C(O)NR''₂, -C(S)NR''₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR'', or -CH₂SR'';

R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R"OR', -R"SR', 25 -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic 30 groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

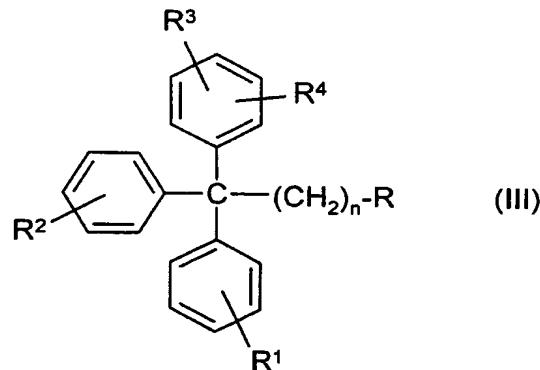
which triaryl methane derivative may further be substituted one or more times with a substituent X selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR", -SR", -R'OR", -R'SR", -C(O)R", -C(S)R", -C(O)OR", -C(S)OR", 5 -C(O)SR", -C(S)SR", -C(O)NR'(OR"), -C(S)NR'(OR"), -C(O)NR'(SR"), -C(S)NR'(SR"), -CH(CN)₂, -C(O)NR"₂, -C(S)NR"₂, -CH[C(O)R"]₂, -CH[C(S)R"]₂, -CH[C(O)OR"]₂, -CH[C(S)OR"]₂, -CH[C(O)SR"]₂, -CH[C(S)SR"]₂, -CH₂OR", or -CH₂SR"; and R' and R", independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

10 In a more preferred embodiment, the triaryl methane derivative for use according to the invention is represented by the general Formula II wherein;

the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

15 the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, 20 thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

In a third preferred embodiment the triaryl methane derivative for use according to the invention is represented by the general Formula III



25 and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5, or 6;

R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R'OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), 30

-C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

R¹, R², R³ and R⁴, independently of each another, represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR", -SR", -R'OR", -R'SR", -C(O)R", -C(S)R", -C(O)OR", -C(S)OR", -C(O)SR", -C(S)SR", -C(O)NR'(OR"), -C(S)NR'(OR"), -C(O)NR'(SR"), -C(S)NR'(SR"), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R"]₂, -CH[C(S)R"]₂, -CH[C(O)OR"]₂, -CH[C(S)OR"]₂, -CH[C(O)SR"]₂, -CH[C(S)SR"]₂, -CH₂OR", or -CH₂SR"; and

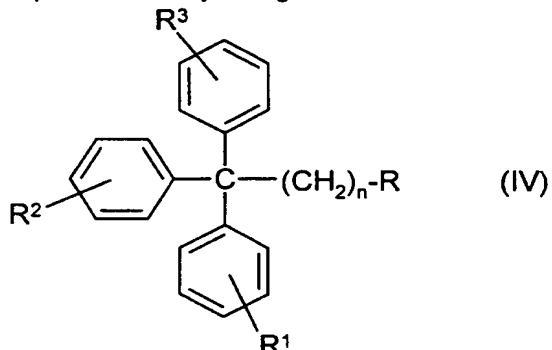
R' and R", independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

In a more preferred embodiment, the triaryl methane derivative for use according to the invention is represented by the general Formula III wherein

the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

In a fourth preferred embodiment the triaryl methane derivative for use according to the invention is represented by the general Formula IV



and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5, or 6;

R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl,

- 5 alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R"OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or
- 10 polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

- 15 R¹, R² and R³, independently of each another, represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR", -SR", -R'OR", -R'SR", -C(O)R", -C(S)R", -C(O)OR", -C(S)OR", -C(O)SR", -C(S)SR", -C(O)NR'(OR"), -C(S)NR'(OR"), -C(O)NR'(SR"), -C(S)NR'(SR"), -CH(CN)₂, -C(O)NR"₂, -C(S)NR"₂, -CH[C(O)R"]₂, -CH[C(S)R"]₂, -CH[C(O)OR"]₂, -CH[C(S)OR"]₂, -CH[C(O)SR"]₂, -CH[C(S)SR"]₂, -CH₂OR", or
- 20 -CH₂SR"; and

R' and R", independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

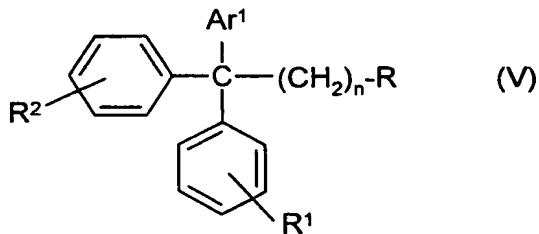
In a more preferred embodiment, the triaryl methane derivative for use according to the invention is represented by the general Formula IV wherein

- 25 the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

In a fifth preferred embodiment the triaryl methane derivative for use according to the invention is represented by the general Formula V

10



and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5, or 6;

5 Ar¹ represents a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'',
10 -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)₂, -C(O)NR''₂, -C(S)NR''₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR'', or -CH₂SR'';

15 R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R'OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or
20 polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

25 R' and R'', independently of each another, represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'', -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)₂, -C(O)NR''₂, -C(S)NR''₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR'', or -CH₂SR''; and

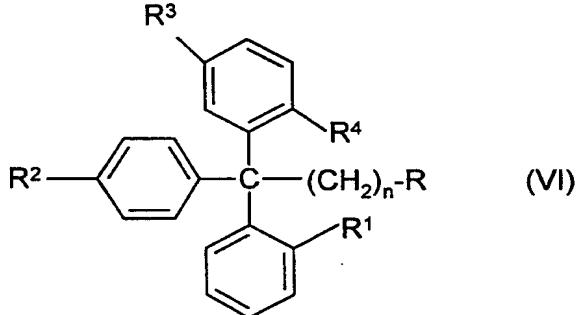
30 R' and R'', independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

In a more preferred embodiment, the triaryl methane derivative for use according to the invention is represented by the general Formula V wherein;

the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic 5 monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

10 In a sixth preferred embodiment the triaryl methane derivative for use according to the invention is represented by the general Formula VI



and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

15 n is 0, 1, 2, 3, 4, 5, or 6;

R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R"OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

130 R^1 , R^2 , R^3 and R^4 , independently of each another, represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR", -SR", -R"OR", -R"SR", -C(O)R", -C(S)R", -C(O)OR", -C(S)OR", -C(O)SR", -C(S)SR", -C(O)NR"(OR"), -C(S)NR"(OR"), -C(O)NR"(SR"), -C(S)NR"(SR"), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R"]₂, -CH[C(S)R"]₂,

-CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; and

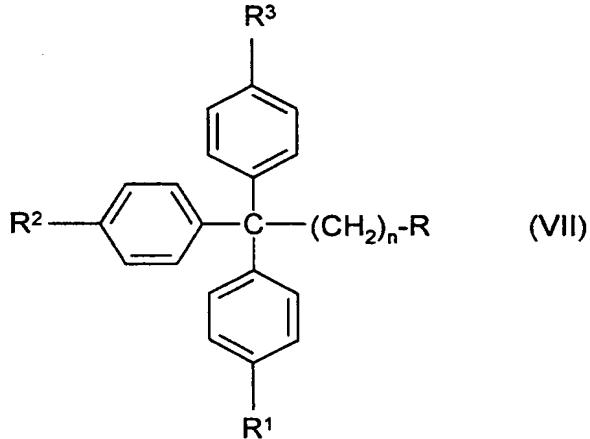
R' and R", independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

5 In a more preferred embodiment, the triaryl methane derivative for use according to the invention is represented by the general Formula VI wherein;

the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

10 the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, 15 thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

In a seventh preferred embodiment the triaryl methane derivative for use according to the invention is represented by the general Formula VII



20 and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5, or 6;

R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R"OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic

groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

R¹, R² and R³, independently of each another, represents hydrogen, 5 halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'', -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)₂, -C(O)NR''₂, -C(S)NR''₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR'', or 10 -CH₂SR''; and

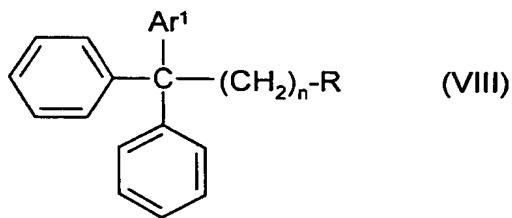
R' and R'', independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

In a more preferred embodiment, the triaryl methane derivative for use according to the invention is represented by the general Formula VII wherein;

15 the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, 20 isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

25 In an eight preferred embodiment the triaryl methane derivative for use according to the invention is represented by the general Formula VIII



and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

30 n is 0, 1, 2, 3, 4, 5, or 6;

Ar¹ represents a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl,

amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'', -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂,

5 -CH₂OR'', or -CH₂SR'';

R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R'OR', -R'SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR'(OR'), -C(S)NR'(OR'), -C(O)NR'(SR'), -C(S)NR'(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

R' and R'', independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

In a more preferred embodiment, the triaryl methane derivative for use according to the invention is represented by the general Formula VIII wherein;

20 the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, 25 isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

30 Definition of Substituents

In the context of this invention halogen represents a fluorine, a chlorine, a bromine or a iodine atom. Thus, a trihalogenmethyl group represents e.g. a trifluoromethyl group and a trichloromethyl group.

In the context of this invention an alkyl group designates a univalent 35 saturated, straight or branched hydrocarbon chain. The hydrocarbon chain preferably contain of from one to eighteen carbon atoms (C₁₋₁₈-alkyl), more preferred a lower alkyl of from one to six carbon atoms (C₁₋₆-alkyl), including pentyl, isopentyl, neopentyl, tertiary pentyl, hexyl and isohexyl. In a preferred embodiment alkyl

represents a C₁₋₄-alkyl group, including butyl, isobutyl, secondary butyl, and tertiary butyl. In a most preferred embodiment alkyl represents a C₁₋₃-alkyl group, which may in particular be methyl, ethyl, propyl or isopropyl.

In the context of this invention a cycloalkyl group designates a cyclic alkyl group, preferably containing of from three to seven carbon atoms (C₃₋₇-cycloalkyl), including cyclopropyl, cyclobutyl, cyclopentyl, and cyclohexyl.

In the context of this invention an alkenyl group designates a carbon chain containing one or more double bonds, including di-enes, tri-enes and poly-enes. In a preferred embodiment the alkenyl group of the invention comprises of from two to six carbon atoms (C₂₋₆-alkenyl), including at least one double bond. In a most preferred embodiment the alkenyl group of the invention is ethenyl; 1,2- or 2,3-propenyl; or 1,2-, 2,3-, or 3,4-butenyl.

In the context of this invention an alkynyl group designates a carbon chain containing one or more triple bonds, including di-ynes, tri-ynes and poly-ynes. In a preferred embodiment the alkynyl group of the invention comprises of from two to six carbon atoms (C₂₋₆-alkynyl), including at least one triple bond. In its most preferred embodiment the alkynyl group of the invention is ethynyl, 1,2- or 2,3-propynyl, 1,2-, 2,3- or 3,4-butynyl.

In the context of this invention an alkoxy group designates an "alkyl-O-" group, wherein alkyl is as defined above.

In the context of this invention an amino group may be a primary (-NH₂), secondary (-NH-alkyl), or tertiary (-N(alkyl)₂) amino group, i.e. it may be substituted once or twice with an alkyl group as defined above.

In the context of this invention a mono- or polycyclic aryl group designates a monocyclic or polycyclic aromatic hydrocarbon group. Examples of preferred aryl groups of the invention are phenyl, biphenyl, naphthyl and anthracenyl.

In the context of this invention a mono- or poly-heterocyclic group is a mono- or polycyclic aromatic group, which holds one or more heteroatoms in its ring structure. Preferred heterocyclic monocyclic groups of the invention are 5- and 6-membered heterocyclic monocyclic groups. Examples of preferred heterocyclic monocyclic groups of the invention include furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolyl, thiadiazolyl, thiazolyl, and thienyl. Examples of preferred heterocyclic polycyclic groups of the invention include benzimidazolyl, indolyl, isoquinolyl and quinolyl.

The chemical compounds for use according to the invention have been described and may be prepared by methods known in the art.

Pharmaceutically Acceptable Salts

The chemical compound for use according to the invention may be provided in any form suitable for the intended administration. Suitable forms include 5 pharmaceutically (i.e. physiologically) acceptable salts, or pre- or prodrug forms of the chemical compound for use according to the invention.

Examples of pharmaceutically acceptable addition salts include, without limitation, the non-toxic inorganic and organic acid addition salts such as the acetate derived from acetic acid, the aconate derived from aconitic acid, the ascorbate 10 derived from ascorbic acid, the benzenesulfonate derived from benzenesulfonic acid, the benzoate derived from benzoic acid, the cinnamate derived from cinnamic acid, the citrate derived from citric acid, the embonate derived from embonic acid, the enantate derived from enanthic acid, the formate derived from formic acid, the fumarate derived from fumaric acid, the glutamate derived from glutamic acid, the glycolate 15 derived from glycolic acid, the hydrochloride derived from hydrochloric acid, the hydrobromide derived from hydrobromic acid, the lactate derived from lactic acid, the maleate derived from maleic acid, the malonate derived from malonic acid, the mandelate derived from mandelic acid, the methanesulfonate derived from methane sulphonic acid, the naphthalene-2-sulphonate derived from naphthalene-2-sulphonic 20 acid, the nitrate derived from nitric acid, the perchlorate derived from perchloric acid, the phosphate derived from phosphoric acid, the phthalate derived from phthalic acid, the salicylate derived from salicylic acid, the sorbate derived from sorbic acid, the stearate derived from stearic acid, the succinate derived from succinic acid, the sulphate derived from sulphuric acid, the tartrate derived from tartaric acid, the 25 toluene-p-sulphonate derived from p-toluene sulphonic acid, and the like. Such salts may be formed by procedures well known and described in the art.

Other acids such as oxalic acid, which may not be considered pharmaceutically acceptable, may be useful in the preparation of salts useful as intermediates in obtaining a chemical compound for use according to the invention 30 and its pharmaceutically acceptable acid addition salt.

Metal salts of a chemical compound for use according to the invention includes alkali metal salts, such as the sodium salt, of a chemical compound for use according to the invention containing a carboxy group.

In the context of this invention the "onium salts" of N-containing compounds 35 are also contemplated as pharmaceutically acceptable salts. Preferred "onium salts" include the alkyl-onium salts, the cycloalkyl-onium salts, and the cycloalkylalkyl-onium salts.

The chemical compound for use according to the invention may be provided in soluble or insoluble forms together with a pharmaceutically acceptable solvents such as water, ethanol, and the like. Soluble forms may also include hydrated forms such as the monohydrate, the dihydrate, the hemihydrate, the 5 trihydrate, the tetrahydrate, and the like. In general, the soluble forms are considered equivalent to insoluble forms for the purposes of this invention.

Steric Isomers

The chemical compounds of the present invention may exist in (+) and (-) 10 forms as well as in racemic forms. The racemates of these isomers and the individual isomers themselves are within the scope of the present invention.

Racemic forms can be resolved into the optical antipodes by known methods and techniques. One way of separating the diastereomeric salts is by use of an optically active acid, and liberating the optically active amine compound by 15 treatment with a base. Another method for resolving racemates into the optical antipodes is based upon chromatography on an optical active matrix. Racemic compounds of the present invention can thus be resolved into their optical antipodes, e.g., by fractional crystallisation of d- or l- (tartrates, mandelates, or camphorsulphonate) salts for example.

20 The chemical compounds of the present invention may also be resolved by the formation of diastereomeric amides by reaction of the chemical compounds of the present invention with an optically active activated carboxylic acid such as that derived from (+) or (-) phenylalanine, (+) or (-) phenylglycine, (+) or (-) camphanic acid or by the formation of diastereomeric carbamates by reaction of the chemical compound of 25 the present invention with an optically active chloroformate or the like.

Additional methods for the resolving the optical isomers are known in the art. Such methods include those described by *Jaques J, Collet A, & Wilen S* in "Enantiomers, Racemates, and Resolutions", John Wiley and Sons, New York (1981).

Moreover, some of the chemical compounds for use according to the 30 invention being oximes, may thus exist in two forms, syn- and anti-form (Z- and E-form), depending on the arrangement of the substituents around the -C=N- double bond. A chemical compound of the present invention may thus be the syn- or the anti-form (Z- and E-form), or it may be a mixture hereof.

35 Biological Activity

As described above, the IK_{Ca} inhibitory compounds for use according to the invention are particularly useful as immune modulating agents, i.e. agents capable of regulating the immune system. More particularly, the IK_{Ca} inhibitory compounds of the

present invention may be used for reducing or inhibiting undesired immune-regulatory actions.

In a preferred embodiment, the invention relates to the use of an IK_{Ca} inhibitory compound for the treatment or alleviation of a diseases, disorders or 5 condition related to immune dysfunction, or in order to obtain immune suppression in an individual in need herefore.

In a more preferred embodiment, the invention relates to the use of an IK_{Ca} inhibitory compound of the invention in a combination therapy with known immune-suppressants for the treatment or alleviation of a diseases, disorders or condition 10 related to immune dysfunction, or for obtaining immune suppression. Preferred immune-suppressants to combine with the compounds of the invention include Amphotericin, Busulphan, Co-trimoxazole, Chlorambucil, colony stimulating factors, corticosteroids, Cyclophosphamide, Fluconazole, folic acid, Ganciclovir, antilymphocyte immunoglobulins, normal immunoglobulins, Methotrexate, 15 Methylprednisolone, Octreotide, Oxpentifyline, Tacrolimus (FK506), Thalidomide, Zolimomab aritox, and the calcineurin inhibitors (protein phosphatase 2B inhibitors), in particular Cyclosporin.

Conditions which may benefit from this treatment include, but are not limited to diseases, disorders or conditions such as auto-immune diseases, e.g. 20 Addison's disease, alopecia areata, Ankylosing spondylitis, haemolytic anemia (anemia haemolytica), pernicious anemia (anemia perniciosa), aphthae, aphthous stomatitis, arthritis, arteriosclerotic disorders, osteoarthritis, rheumatoid arthritis, aspermiogenesis, asthma bronchiale, auto-immune asthma, auto-immune hemolysis, Bechet's disease, Boeck's disease, inflammatory bowel disease, Burkitt's lymphoma, 25 Chron's disease, chorioiditis, colitis ulcerosa, Coeliac disease, cryoglobulinemia, dermatitis herpetiformis, dermatomyositis, insulin-dependent type I diabetes, juvenile diabetes, idiopathic diabetes insipidus, insulin-dependent diabetes mellitis, auto-immune demyelinating diseases, Dupuytren's contracture, encephalomyelitis, encephalomyelitis allergica, endophthalmia phacoanaphylactica, enteritis allergica, 30 autoimmune enteropathy syndrome, erythema nodosum leprosum, idiopathic facial paralysis, chronic fatigue syndrome, febris rheumatica, glomerulo nephritis, Goodpasture's syndrome, Graves' disease, Hamman-Rich's disease, Hashimoto's disease, Hashimoto's thyroiditis, sudden hearing loss, sensoneural hearing loss, hepatitis chronica, Hodgkin's disease, haemoglobinuria paroxysmata, 35 hypogonadism, ileitis regionalis, iritis, leucopenia, leucemia, lupus erythematosus disseminatus, systemic lupus erythematosus, cutaneous lupus erythematosus, lymphogranuloma malignum, mononucleosis infectiosa, myasthenia gravis, traverse myelitis, primary idiopathic myxedema, nephrosis, ophthalmia sympathica, orchitis

granulomatosa, pancreatitis, pemphigus, pemphigus vulgaris, polyarteritis nodosa, polyarthritis chronica primaria, polymyositis, polyradiculitis acuta, psoriasis, purpura, pyoderma gangrenosum, Quervain's thyreoiditis, Reiter's syndrome, sarcoidosis, ataxic sclerosis, progressive systemic sclerosis, scleritis, sclerodermia, multiple 5 sclerosis, sclerosis disseminata, acquired spenic atrophy, infertility due to antispermatozoan antibodies, thrombocytopenia, idiopathic thrombocytopenia purpura, thymoma, acute anterior uveitis, vitiligo, AIDS, HIV, SCID and Epstein Barr virus associated diseases such as Sjogren's syndrome, virus (AIDS or EBV) associated B cell lymphoma, parasitic diseases such as Leishmania, and immune- 10 suppressed disease states such as viral infections following allograft transplantations, graft vs. Host syndrome, transplant rejection, or AIDS, cancer, chronic active hepatitis diabetes, toxic shock syndrome, food poisoning, and transplant rejection.

Accordingly, in further embodiments, the invention relates to a chemical compound having IK_{Ca} inhibitory activity for use as a medicament.

15 More specifically the invention relates to the use of a chemical compound having selective IK_{Ca} inhibitory activity for use in the manufacture of a medicament for the treatment of treatment of diseases related to immune dysfunction. In a preferred embodiment the medicament is an immune system suppressing medicament (an immune-suppressivum).

20

Pharmaceutical Compositions

In yet another aspect the invention relates to pharmaceutical compositions for use in the treatment or alleviation of diseases, disorders or conditions related to immune dysfunction, which pharmaceutical composition comprises a therapeutically 25 effective amount of a chemical compound having IK_{Ca} inhibitory activity, as identified by the method of the invention.

While a chemical compound for use according to the invention for use in therapy may be administered in the form of the raw chemical compound, it is preferred to introduce the active ingredient, optionally in the form of a physiologically 30 acceptable salt, in a pharmaceutical composition together with one or more adjuvants, excipients, carriers, buffers, diluents, and/or other customary pharmaceutical auxiliaries.

In a preferred embodiment, the invention provides pharmaceutical compositions comprising the chemical compound for use according to the invention or 35 a pharmaceutically acceptable salt or derivative thereof together with one or more pharmaceutically acceptable carriers therefor and, optionally, other therapeutic and/or prophylactic ingredients. The carrier(s) must be "acceptable" in the sense of being

compatible with the other ingredients of the formulation and not deleterious to the recipient thereof.

The pharmaceutical composition of the invention may be administered by any convenient route which suite the desired therapy. Preferred routes of 5 administration include oral administration, in particular in tablet, in capsule, in dragé, in powder, or in liquid form, and parenteral administration, in particular cutaneous, subcutaneous, intramuscular, or intravenous injection. The pharmaceutical composition may be prepared by the skilled person using standard and conventional techniques appropriate to the desired formulation. When desired, compositions 10 adapted to give sustained release of the active ingredient may be employed.

The actual dosage depend on the nature and severity of the disease being treated, and is within the discretion of the physician, and may be varied by titration of the dosage to the particular circumstances of this invention to produce the desired therapeutic effect. However, it is presently contemplated that pharmaceutical 15 compositions containing of from about 0.1 to about 500 mg of active ingredient per individual dose, preferably of from about 1 to about 100 mg, most preferred of from about 1 to about 10 mg, are suitable for therapeutic treatments.

Further details on techniques for formulation and administration may be found in the latest edition of Remington's Pharmaceutical Sciences (Maack Publishing 20 Co., Easton, PA).

The actual dosage depend on the nature and severity of the disease being treated, and is within the discretion of the physician, and may be varied by titration of the dosage to the particular circumstances of this invention to produce the desired therapeutic effect. However, it is presently contemplated that pharmaceutical 25 compositions containing of from about 0.1 to about 500 mg of active ingredient per individual dose, preferably of from about 1 to about 100 mg, most preferred of from about 1 to about 10 mg, are suitable for therapeutic treatments.

The active ingredient may be administered in one or several doses per day. A satisfactory result can, in certain instances, be obtained at a dosage as low as 0.1 30 $\mu\text{g}/\text{kg}$ i.v. and $1 \mu\text{g}/\text{kg}$ p.o. The upper limit of the dosage range is presently considered to be about $10 \text{ mg}/\text{kg}$ i.v. and $100 \text{ mg}/\text{kg}$ p.o. Preferred ranges are from about $0.1 \mu\text{g}/\text{kg}$ to about $10 \text{ mg}/\text{kg}/\text{day}$ i.v., and from about $1 \mu\text{g}/\text{kg}$ to about $100 \text{ mg}/\text{kg}/\text{day}$ p.o.

Methods of Therapy

35 Viewed from another aspect, the invention provides a method for treatment, prevention or alleviation of a disease or a disorder or a condition of a living animal body, including a human, which disease, disorder or condition is responsive to IK_{Ca} inhibitory compounds.

Therefore, in a preferred embodiment, the invention provides a method of treatment or alleviation of diseases, disorders or conditions relating to immune dysfunction in a living body, said method comprising administering to said living body an effective amount of a chemical compound having IK_{Ca} inhibitory activity.

5 In a more preferred embodiment, the disease, disorder or condition relating to immune dysfunction is an auto-immune disease, e.g. Addison's disease, alopecia areata, Ankylosing spondylitis, haemolytic anemia (anemia haemolytica), pernicious anemia (anemia perniciosa), aphthae, aphthous stomatitis, arthritis, arteriosclerotic disorders, osteoarthritis, rheumatoid arthritis, aspermiogenese, asthma bronchiale,
10 auto-immune asthma, auto-immune hemolysis, Bechet's disease, Boeck's disease, inflammatory bowel disease, Burkitt's lymphoma, Chron's disease, chorioiditis, colitis ulcerosa, Coeliac disease, cryoglobulinemia, dermatitis herpetiformis, dermatomyositis, insulin-dependent type I diabetes, juvenile diabetes, idiopathic diabetes insipidus, insulin-dependent diabetes mellitis, auto-immune demyelinating
15 diseases, Dupuytren's contracture, encephalomyelitis, encephalomyelitis allergica, endophthalmia phacoanaphylactica, enteritis allergica, autoimmune enteropathy syndrome, erythema nodosum leprosum, idiopathic facial paralysis, chronic fatigue syndrome, febris rheumatica, glomerulo nephritis, Goodpasture's syndrome, Graves' disease, Hamman-Rich's disease, Hashimoto's disease, Hashimoto's thyroiditis,
20 sudden hearing loss, sensoneural hearing loss, hepatitis chronica, Hodgkin's disease, haemoglobinuria paroxysmatica, hypogonadism, ileitis regionalis, iritis, leucopenia, leucemia, lupus erythematosus disseminatus, systemic lupus erythematosus, cutaneous lupus erythematosus, lymphogranuloma malignum, mononucleosis infectiosa, myasthenia gravis, traverse myelitis, primary idiopathic myxedema,
25 nephrosis, ophthalmia sympathica, orchitis granulomatosa, pancreatitis, pemphigus, pemphigus vulgaris, polyarteritis nodosa, polyarthritis chronica primaria, polymyositis, polyradiculitis acuta, psoriasis, purpura, pyoderma gangrenosum, Quervain's thyroiditis, Reiter's syndrome, sarcoidosis, ataxic sclerosis, progressive systemic sclerosis, scleritis, sclerodermia, multiple sclerosis, sclerosis disseminata, acquired
30 spenic atrophy, infertility due to antispermatozoan antibodies, thrombocytopenia, idiopathic thrombocytopenia purpura, thymoma, acute anterior uveitis, vitiligo, AIDS, HIV, SCID and Epstein Barr virus associated diseases such as Sjogren's syndrome, virus (AIDS or EBV) associated B cell lymphoma, parasitic diseases such as Leslihmania, and immune-suppressed disease states such as viral infections following
35 allograft transplantations, graft vs. Host syndrome, transplant rejection, or AIDS, cancer, chronic active hepatitis diabetes, toxic chock syndrome, food poisoning, or transplant rejection.

In another preferred embodiment, the method of the invention comprises simultaneous administration of the chemical compound having selective IK_{Ca} inhibitory activity and a pharmaceutically effective amount of a conventional immune suppressing agent.

5 In a more preferred embodiment the immune-suppressing agent is Amphotericin, Busulphan, Co-trimoxazole, Chlorambucil, colony stimulating factors, corticosteroids, Cyclophosphamide, Fluconazole, folinic acid, Ganciclovir, antilymphocyte immunoglobulins, normal immunoglobulins, Methotrexate, Methylprednisolone, Octreotide, Oxpentifyline, Tacrolimus (FK506), Thalidomide, 10 Zolimomab aritox, or the calcineurin inhibitors (protein phosphatase 2B inhibitors), in particular Cyclosporin.

It is at present contemplated that suitable dosage ranges are 0.1 to 1000 milligrams daily, 10-500 milligrams daily, and especially 30-100 milligrams daily, dependent as usual upon the exact mode of administration, form in which 15 administered, the indication toward which the administration is directed, the subject involved and the body weight of the subject involved, and further the preference and experience of the physician or veterinarian in charge.

BRIEF DESCRIPTION OF THE DRAWINGS

20

The present invention is further illustrated by reference to the accompanying drawing, in which:

Fig. 1 shows the effect of a compound of the invention (Clotrimazole) on Cyclosporin A mediated inhibition of T cell proliferation (PPD-induced T cell 25 proliferation) on a relative scale of from 0,00 to 1,25, carried out as described in Example 2 [with and without (Control) Clotrimazole, 10 μ M; Combined with Cyclosporin A, in concentrations of 0, 2.5, 5, 10 and 25 nM, respectively].

EXAMPLES

30

The invention is further illustrated with reference to the following examples which are not intended to be in any way limiting to the scope of the invention as claimed.

35 **Example 1**

Inhibition of T Cell Proliferation

The chemical compounds used according to the invention prevent immunological proliferation by selective inhibition of the Ca^{2+} -activated K-channels in

T- and B-lymphocytes. This effect may be verified using various proliferation assays. In this experiment the proliferative assay described by Ødum *et al.* [Ødum N, Kanner S B, Ledbetter J A, & Svejgaard A; *J. Immunol.* 1993 **150** (12) 5289-5298] was used.

5 The chemical compounds representative for the invention tested in this experiment are (4-chlorophenyl-diphenyl)-carbinol (A); ethyl-2-phenyl-2-(1-piperidyl)-phenylacetate (B); and 1,1,1-triphenylacetone (C); all compounds commercially available from Sigma-Aldrich, Denmark.

Assays were performed in culture medium (RPMI 1640; available from Gibco, Grand Island, NY) supplemented with 10% pooled human serum, 2 mM L-10 glutamine, 100 µg/ml penicillin, and 100 µg/ml streptomycin (available from Novo Nordisk, Copenhagen, Denmark) in 96-well round bottom tissue culture plates (available from Nunc, Roskilde, Denmark) with a final volume of 200 µl.

T cells were pre-incubated for three hours with the test compounds before addition of antigen (PPD; Purified protein derivative, available from Statens Serum Institut, Denmark; 100 µg/ml). T cells were cultured at 5×10^4 cells/well for 144 hours. 15 Twelve hours before harvest, [³H]thymidine (1 x Ci/well) was added. The cells were harvested onto glass fibre filters, and the [³H]thymidine incorporation was measured in a scintillation counter. The results were expressed as mean counts per minute (cpm) from triplicate cultures.

20 The results are presented in Table 1, below.

Table 1
Inhibition of T Cell Proliferation

Test Compound	Medium Solvent	Solvent	T Cell Proliferation (cpm x 10 ⁻³)		
			Antigen, PPD		
			2.5 µM	10 µM	25 µM
A	0.2	26.1	21.5	19.8	18.1
B	0.2	26.1	22.5	20	19
C	0.2	26.1	25.5	18	19

25 These results show that the number of T cells decreases in the presence of increasing concentrations of the chemical compound for use according to the invention, and support the fact that the chemical compounds for use according to the invention inhibit the antigen induced T cell proliferation and thus are useful for the reduction or inhibition of undesired immune-regulatory actions.

Example 2**Combination Treatment**

In this example, the effect of a compound of this invention (Clotrimazole) on Cyclosporin A mediated inhibition of T cell proliferation is determined.

5 T cells were stimulated with antigen in the presence of Cyclosporin A, or Cyclosporin A and Clotrimazole, respectively.

The proliferation assay described in Example 1, was used.

Cells were incubated for 5 days in culture medium with *PPD* in the presence of Cyclosporin A, or Cyclosporin A and Clotrimazole, respectively.

10 Clotrimazole (10 μ M) was added 30 minutes prior to the addition of antigen. [3 H]thymidine (1 mM Ci) incorporation was then measured in triplicate wells. The bars shown in Fig. 1 represent 3 independent experiments \pm S.E. ($p \leq 0.05$ vs. control). Eleven other experiments using *Candida albicans* antigen, *tetanus toxin*, Con A or PHA as the antigen/mitogen challenge gave similar results.

15 T cell proliferation was assayed 6 days after stimulation using 3 H-thymidine incorporation. The Cyclosporin A mediated inhibition of T cell proliferation is shifted leftwards by 10 μ M Clotrimazole, from a 50% inhibition of proliferation at approximately 25 nM Cyclosporin A to half-maximal inhibition at 2.5 nM Cyclosporin A.

20 This suggests that the antigen-induced T cell proliferation is highly sensitive to both IK channel block and inhibition of calcineurin, and data indicate that the IK channel is highly important for normal T cell proliferation and suggest that IK channels are attractive targets for immune suppression.

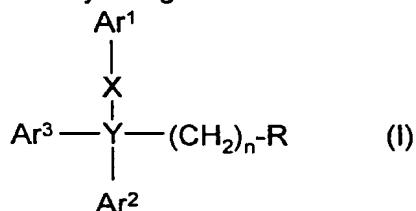
CLAIMS:

1. Use of a chemical compound having selective IK_{Ca} modulatory activity for the manufacture of a medicament for the treatment, prevention or alleviation of a disease or a disorder or a condition of a mammal, including a human, which disease, disorder or condition relates to immune dysfunction.

5

2. The use according to claim 1, wherein the chemical compound is a triaryl methane derivative represented by the general Formula I

10



and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

15 n is 0, 1, 2, 3, 4, 5 or 6;

20 X is absent, or represent a group of the formula $-(\text{CH}_2)_n-$, of the formula $-(\text{CH}_2)_n-\text{Z}-$ (in either direction), of the formula $-(\text{CH}_2)_n-\text{CH}=\text{N}-$ (in either direction), the formula $-(\text{CH}_2)_n-\text{Z}-(\text{CH}_2)_m-$, or of the formula $-(\text{CH}_2)_n-\text{CH}=\text{N}-(\text{CH}_2)_m-$ (in either direction), or a group of the formula $-\text{R}''' \text{C}(\text{O})\text{N}-$;

in which formulas

n and m , independently of each another, represent 0, 1, 2, 3 or 4; and Z represents O, S, or NR'' , wherein R'' represents hydrogen or alkyl;

25 Y represents a carbon atom (C), a nitrogen atom (N), or a phosphor atom (P), a silicium atom (Si), or a germanium atom (Ge);

30 Ar^1 , Ar^2 and Ar^3 , independently of each another, represents a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-
heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula $-\text{OR}''$, $-\text{SR}''$, $-\text{R}'\text{OR}''$, $-\text{R}'\text{SR}''$, $-\text{C}(\text{O})\text{R}''$, $-\text{C}(\text{S})\text{R}''$, $-\text{C}(\text{O})\text{OR}''$, $-\text{C}(\text{S})\text{OR}''$, $-\text{C}(\text{O})\text{SR}''$, $-\text{C}(\text{S})\text{SR}''$, $-\text{C}(\text{O})\text{NR}'(\text{OR}')$, $-\text{C}(\text{S})\text{NR}'(\text{OR}')$, $-\text{C}(\text{O})\text{NR}'(\text{SR}')$, $-\text{C}(\text{S})\text{NR}'(\text{SR}')$, $-\text{CH}(\text{CN})_2$, $-\text{C}(\text{O})\text{NR}''_2$,

35

-C(S)NR"₂, -CH[C(O)R"]₂, -CH[C(S)R"]₂, -CH[C(O)OR"]₂, -CH[C(S)OR"]₂, -CH[C(O)SR"]₂, -CH[C(S)SR"]₂, -CH₂OR", or -CH₂SR";

R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R"OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR'; and

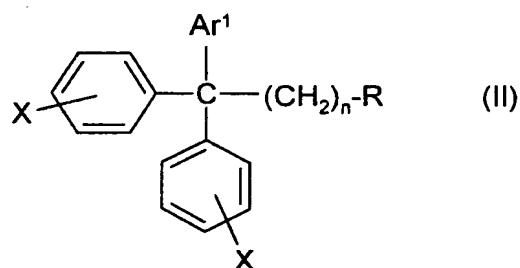
15 R' and R", independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

3. The use according to claim 2, wherein

20 the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene;

25 and the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

30 4. The use according to claim 2, wherein the chemical compound is a triaryl methane derivative represented by the general Formula II



and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5 or 6;

5 Ar¹ represents a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR", -SR", -R'OR", -R'SR", -C(O)R", -C(S)R", -C(O)OR", -C(S)OR", -C(O)SR", -C(S)SR", -C(O)NR'(OR"), -C(S)NR'(OR"), -C(O)NR'(SR"), -C(S)NR'(SR"), -CH(CN)₂, -C(O)NR"₂, -C(S)NR"₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR", or -CH₂SR";

10 R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R"OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; or a 15 partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

20 which triaryl methane derivative may further be substituted one or more times with a substituent X selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR", -SR", -R'OR", -R'SR", -C(O)R", -C(S)R", -C(O)OR", -C(S)OR", -C(O)SR", -C(S)SR", -C(O)NR'(OR"), -C(S)NR'(OR"), -C(O)NR'(SR"), -C(S)NR'(SR"), -CH(CN)₂, -C(O)NR"₂, -C(S)NR"₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR", or -CH₂SR"; and

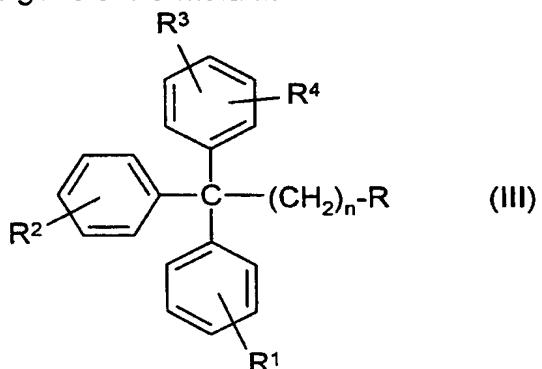
25 R' and R", independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

30 35 5. The use according to claim 4, wherein

the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, 5 pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

6. The use according to claim 2, wherein the triaryl methane derivative is represented by the general Formula III



10

and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5, or 6;

15 R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R"OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)2, -C(O)NR'2, -C(S)NR'2, -CH[C(O)R']2, -CH[C(S)R']2, -CH[C(O)OR']2, -CH[C(S)OR']2, -CH[C(O)SR']2, -CH[C(S)SR']2, -CH2OR', or -CH2SR'; or a 20 partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

25 R¹, R², R³ and R⁴, independently of each another, represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'', -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)2, -C(O)NR'2, -C(S)NR'2, -CH[C(O)R'']2,

-CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR'', or -CH₂SR''; and

R' and R'', independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

5

7. The use according to claim 6, wherein

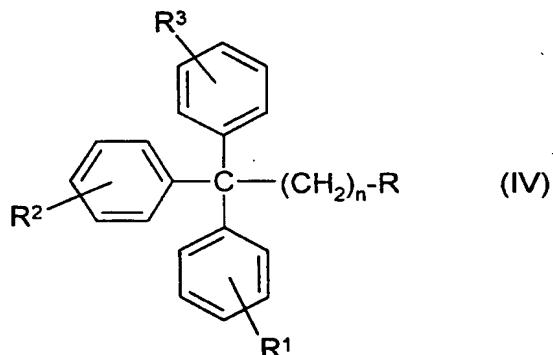
the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

10

the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

15

8. The use according to claim 2, wherein the triaryl methane derivative is represented by the general Formula IV



20

and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5, or 6;

25

R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R''OR', -R''SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR''(OR'), -C(S)NR''(OR'), -C(O)NR''(SR'), -C(S)NR''(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or polycyclic aryl group, or a mono- or

30

poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

5 R¹, R² and R³, independently of each another, represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'', -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)₂, -C(O)NR''₂, -C(S)NR''₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR'', or -CH₂SR''; and

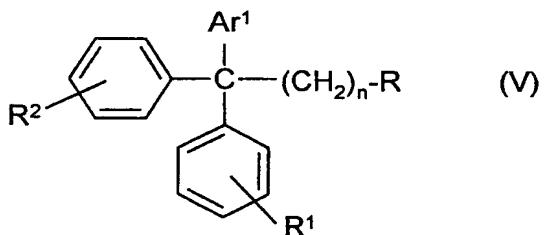
10 R' and R'', independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

15 9. The use according to claim 8, wherein

the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

20 the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

25 10. The use according to claim 2, wherein the triaryl methane derivative is represented by the general Formula V



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and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5, or 6;

5 Ar¹ represents a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'', -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)₂, -C(O)NR''₂, -C(S)NR''₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR'', or -CH₂SR'';

10 R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R"OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

15 R¹ and R², independently of each another, represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'', -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)₂, -C(O)NR''₂, -C(S)NR''₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR'', or -CH₂SR''; and

20 R' and R'', independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

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11. The use according to claim 10, wherein

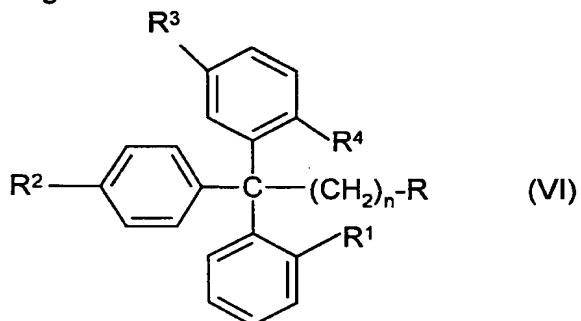
the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

35

the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl,

piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

5 12. The use according to claim 2, wherein the triaryl methane derivative is represented by the general Formula VI



and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

10 n is 0, 1, 2, 3, 4, 5, or 6;

R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R"OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

25 R¹, R², R³ and R⁴, independently of each another, represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'', -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR'', or -CH₂SR''; and

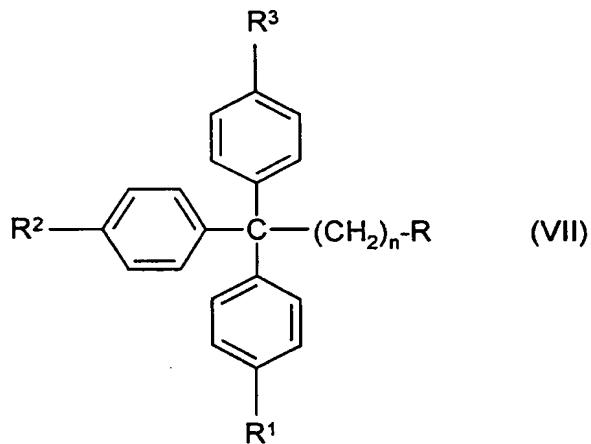
30 R' and R'', independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

13. The use according to claim 12, wherein

the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

5 the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, 10 pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

14. The use according to claim 2, wherein the triaryl methane derivative is represented by the general Formula VII



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and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5, or 6;

20 R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R"OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)2, -C(O)NR'2, -C(S)NR'2, -CH[C(O)R']2, -CH[C(S)R']2, -CH[C(O)OR']2, -CH[C(S)OR']2, -CH[C(O)SR']2, -CH[C(S)SR']2, -CH₂OR', or -CH₂SR'; or a 25 partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group

consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

5 R¹, R² and R³, independently of each another, represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'', -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)₂, -C(O)NR''₂, -C(S)NR''₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR'', or -CH₂SR''; and

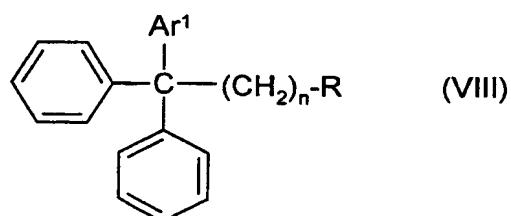
10 R' and R'', independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

15. The use according to claim 14, wherein

15 the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

20 the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

25 16. The use according to claim 2, wherein the triaryl methane derivative is represented by the general Formula VIII



30 and a pharmaceutically acceptable salt or an oxide or a hydrate thereof, wherein,

n is 0, 1, 2, 3, 4, 5, or 6;

Ar¹ represents a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from

the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR'', -SR'', -R'OR'', -R'SR'', -C(O)R'', -C(S)R'', -C(O)OR'', -C(S)OR'', -C(O)SR'', -C(S)SR'', -C(O)NR'(OR''), -C(S)NR'(OR''), -C(O)NR'(SR''), -C(S)NR'(SR''), -CH(CN)₂, -C(O)NR''₂, -C(S)NR''₂, -CH[C(O)R'']₂, -CH[C(S)R'']₂, -CH[C(O)OR'']₂, -CH[C(S)OR'']₂, -CH[C(O)SR'']₂, -CH[C(S)SR'']₂, -CH₂OR'', or -CH₂SR'';

5 R represents hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', -SR', -R"OR', -R"SR', -C(O)R', -C(S)R', -C(O)OR', -C(S)OR', -C(O)SR', -C(S)SR', -C(O)NR"(OR'), -C(S)NR"(OR'), -C(O)NR"(SR'), -C(S)NR"(SR'), -CH(CN)₂, -C(O)NR'₂, -C(S)NR'₂, -CH[C(O)R']₂, -CH[C(S)R']₂, -CH[C(O)OR']₂, -CH[C(S)OR']₂, -CH[C(O)SR']₂, -CH[C(S)SR']₂, -CH₂OR', or -CH₂SR'; or a partially or completely saturated mono- or polycyclic aryl group, or a mono- or poly-heterocyclic group, which mono- or polycyclic groups may optionally be substituted one or more times with substituents selected from the group consisting of hydrogen, halogen, trihalogenmethyl, alkyl, cycloalkyl, alkenyl, alkynyl, amino, nitro or cyano, or a group of the formula -OR', or -SR';

10 R' and R'', independently of each another, represents hydrogen, alkyl, cycloalkyl, alkenyl, alkynyl, or alkoxy.

20

17. The use according to claim 16, wherein

the partially or completely saturated mono- or polycyclic aryl group is selected from the group consisting phenyl, biphenyl, naphthyl, or cyclopenta-2,4-diene-1-ylidene; and

25

the mono- or poly-heterocyclic group is A 5- and 6 membered heterocyclic monocyclic group selected from the group consisting of furanyl, imidazolyl, isoimidazolyl, 2-isoimidazolyl, isothiazolyl, isoxazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolyl, piperidyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolidinyl, 30 pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and butyrolactonyl, in particular γ -butyrolactonyl.

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18. The use according to claim 2, wherein the compound is

(4-chlorophenyl-diphenyl)-carbinol;

Ethyl 2-phenyl-2-(1-piperidyl)-phenylacetate; or

1,1,1-triphenylacetone;

or a pharmaceutically acceptable salt or an oxide or a hydrate hereof.

19. The use according to any of claims 1-18, wherein the disease, disorder or condition relating to immune dysfunction is an auto-immune disease, e.g. Addison's disease, alopecia areata, Ankylosing spondylitis, haemolytic anemia (anemia haemolytica), pernicious anemia (anemia perniciosa), aphthae, 5 aphthous stomatitis, arthritis, arteriosclerotic disorders, osteoarthritis, rheumatoid arthritis, aspermiogenesis, asthma bronchiale, auto-immune asthma, auto-immune hemolysis, Bechet's disease, Boeck's disease, inflammatory bowel disease, Burkitt's lymphoma, Chron's disease, chorioiditis, colitis ulcerosa, Coeliac disease, cryoglobulinemia, dermatitis herpetiformis, dermatomyositis, 10 insulin-dependent type I diabetes, juvenile diabetes, idiopathic diabetes insipidus, insulin-dependent diabetes mellitus, auto-immune demyelinating diseases, Dupuytren's contracture, encephalomyelitis, encephalomyelitis allergica, endophthalmia phacoanaphylactica, enteritis allergica, autoimmune enteropathy syndrome, erythema nodosum leprosum, idiopathic facial paralysis, 15 chronic fatigue syndrome, febris rheumatica, glomerulo nephritis, Goodpasture's syndrome, Graves' disease, Hamman-Rich's disease, Hashimoto's disease, Hashimoto's thyroiditis, sudden hearing loss, sensoneural hearing loss, hepatitis chronica, Hodgkin's disease, haemoglobinuria paroxysmata, hypogonadism, ileitis regionalis, iritis, leucopenia, leucemia, lupus erythematosus disseminatus, 20 systemic lupus erythematosus, cutaneous lupus erythematosus, lymphogranuloma malignum, mononucleosis infectiosa, myasthenia gravis, traverse myelitis, primary idiopathic myxedema, nephrosis, ophthalmia sympathica, orchitis granulomatosa, pancreatitis, pemphigus, pemphigus vulgaris, polyarteritis nodosa, polyarthritis chronica primaria, polymyositis, 25 polyradiculitis acuta, psoriasis, purpura, pyoderma gangrenosum, Quervain's thyroiditis, Reiter's syndrome, sarcoidosis, ataxic sclerosis, progressive systemic sclerosis, scleritis, sclerodermia, multiple sclerosis, sclerosis disseminata, acquired spenic atrophy, infertility due to antispermatozoan antibodies, thrombocytopenia, idiopathic thrombocytopenia purpura, thymoma, 30 acute anterior uveitis, vitiligo, AIDS, HIV, SCID and Epstein Barr virus associated diseases such as Sjorgren's syndrome, virus (AIDS or EBV) associated B cell lymphoma, parasitic diseases such as Leishmania, and immune-suppressed disease states such as viral infections following allograft transplantations, graft vs. Host syndrome, transplant rejection, or AIDS, cancer, chronic active hepatitis 35 diabetes, toxic shock syndrome, food poisoning, or transplant rejection.

20. The use according to claims 1-19, for the manufacture of a medicament which medicament further comprises a pharmaceutically effective amount of a conventional immune suppressing agent.
- 5 21. The use according to claim 20, wherein the immune-suppressing agent is Amphotericin, Busulphan, Co-trimoxazole, Chlorambucil, colony stimulating factors, corticosteroids, Cyclophosphamide, Fluconazole, folinic acid, Ganciclovir, antilymphocyte immunoglobulins, normal immunoglobulins, Methotrexate, Methylprednisolone, Octreotide, Oxpentifyline, Tacrolimus (FK506), Thalidomide, Zolimomab aritox, or the calcineurin inhibitors (protein phosphatase 2B inhibitors), in particular Cyclosporin.
- 10 22. A method for of treatment, prevention or alleviation of a disease or a disorder or a condition related to immune dysfunction, which method comprises the step of administering to such a living animal body in need thereof a therapeutically effective amount of a chemical compound having selective IK_{Ca} inhibitory activity.
- 15 23. The method according to claim 22, wherein the disease, disorder or condition relating to immune dysfunction is an auto-immune disease, e.g. Addison's disease, alopecia areata, Ankylosing spondylitis, haemolytic anemia (anemia haemolytica), pernicious anemia (anemia perniciosa), aphthae, aphthous stomatitis, arthritis, arteriosclerotic disorders, osteoarthritis, rheumatoid arthritis, aspermiogenesis, asthma bronchiale, auto-immune asthma, auto-immune 20 hemolysis, Bechet's disease, Boeck's disease, inflammatory bowel disease, Burkitt's lymphoma, Chron's disease, chorioiditis, colitis ulcerosa, Coeliac disease, cryoglobulinemia, dermatitis herpetiformis, dermatomyositis, insulin-dependent type I diabetes, juvenile diabetes, idiopathic diabetes insipidus, insulin-dependent diabetes mellitus, auto-immune demyelinating diseases, 25 Dupuytren's contracture, encephalomyelitis, encephalomyelitis allergica, endophthalmia phacoanaphylactica, enteritis allergica, autoimmune enteropathy syndrome, erythema nodosum leprosum, idiopathic facial paralysis, chronic fatigue syndrome, febris rheumatica, glomerulo nephritis, Goodpasture's syndrome, Graves' disease, Hamman-Rich's disease, Hashimoto's disease, 30 Hashimoto's thyroiditis, sudden hearing loss, sensoneural hearing loss, hepatitis chronica, Hodgkin's disease, haemoglobinuria paroxysmata, hypogonadism, ileitis regionalis, iritis, leucopenia, leucemia, lupus erythematosus disseminatus, systemic lupus erythematosus, cutaneous lupus erythematosus,

lymphogranuloma malignum, mononucleosis infectiosa, myasthenia gravis, traverse myelitis, primary idiopathic myxedema, nephrosis, ophthalmia sympathica, orchitis granulomatosa, pancreatitis, pemphigus, pemphigus vulgaris, polyarteritis nodosa, polyarthritis chronica primaria, polymyositis, 5 polyradiculitis acuta, psoriasis, purpura, pyoderma gangrenosum, Quervain's thyreoiditis, Reiter's syndrome, sarcoidosis, ataxic sclerosis, progressive systemic sclerosis, scleritis, sclerodermia, multiple sclerosis, sclerosis disseminata, acquired spenic atrophy, infertility due to antispermatozoan antibodies, thrombocytopenia, idiopathic thrombocytopenia purpura, thymoma, 10 acute anterior uveitis, vitiligo, AIDS, HIV, SCID and Epstein Barr virus associated diseases such as Sjorgren's syndrome, virus (AIDS or EBV) associated B cell lymphoma, parasitic diseases such as Leishmania, and immune-suppressed disease states such as viral infections following allograft transplantations, graft vs. Host syndrome, transplant rejection, or AIDS, cancer, chronic active hepatitis, 15 diabetes, toxic shock syndrome, food poisoning, or transplant rejection.

24. The method according to either of claims 22-23, which method comprises simultaneous administration of the chemical compound having selective IK_{Ca} inhibitory activity and a pharmaceutically effective amount of a conventional immune suppressing agent.

25. The method according to claim 24, wherein the immune-suppressing agent is Amphotericin, Busulphan, Co-trimoxazole, Chlorambucil, colony stimulating factors, corticosteroids, Cyclophosphamide, Fluconazole, folinic acid, Ganciclovir, antilymphocyte immunoglobulins, normal immunoglobulins, Methotrexate, Methylprednisolone, Octreotide, Oxpentifylline, Tacrolimus (FK506), Thalidomide, Zolimomab artox, or the calcineurin inhibitors (protein phosphatase 2B inhibitors), in particular Cyclosporin.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/DK 00/00253

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: A61K 31/4453, A61K 31/12, A61K 31/055, A61K 31/00, A61P 37/00
According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	WO 9925347 A2 (NEUROSEARCH A/S), 27 May 1999 (27.05.99), claims 1, 17-23; page 1, line 1 - page 5, line 20; page 11, line 14 - line 23; page 20, line 1 - page 21, line 15 --	1-25
X	Cell Calcium, Volume 21, No 1, 1997, Jos A.H. Verheugen et al, "Enhancement of calcium signaling and proliferation responses in activated human T lymphocytes", page 1 - page 17, the abstract; introduction page 16, ultimate paragraph --	1,22

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents:	
"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier document but published on or after the international filing date	"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
6 Sept 2000	28.09.2000
Name and mailing address of the International Searching Authority European Patent Office P.B. 5818 Patentaan 2 NL-2280 HV Rijswijk Tel(+31-70)340-2040, Tx 31 651 epo nl, Fax(+31-70)340-3016	Authorized officer GERD STRANDELL/EÖ Telephone No.

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INTERNATIONAL SEARCH REPORTInternational application No.
PCT/DK 00/00253

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	The Journal of Immunology, Volume 156, 1996, Randall K. Rader et al, "T Cell Activation Is Regulated by Voltage-Dependent and Calcium-Activated Potassium Channels", page 1425 - page 1430, the abstract; figure 4 --	1,22
X	WO 9734589 A1 (PRESIDENT AND FELLOWS OF HARVARD COLLEGE), 25 Sept 1997 (25.09.97), page 13, line 1 - page 19, line 26; the claims, page 27, no. 37; page 28, no. 43; page 33, no. 87 --	1-25
X	WO 9418967 A1 (PRESIDENT AND FELLOWS OF HARVARD COLLEGE), 1 Sept 1994 (01.09.94), page 12, last paragraph . page 13; page 5, third paragraph - page 6; the claims --	1-25
A	US 5540931 A (CHARLES W. HEWITT ET AL), 30 July 1996 (30.07.96) --	20,21,24,25
A	WO 9601107 A1 (HOFMANN, BO, ARNE), 18 January 1996 (18.01.96), claims 1-13 -- -----	1-25

INTERNATIONAL SEARCH REPORTInternational application No.
PCT/DK00/00253**Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)**

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: **22-25**
because they relate to subject matter not required to be searched by this Authority, namely:
see next sheet
2. Claims Nos.: **1-17, 19-25 all in part**
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).:

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.
 No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/DK00/00253

Box I.1

Claims 22-25 relate to methods of treatment of the human or animal body by surgery or by therapy/ diagnostic methods practised on the human or animal body/Rule 39.1.(iv). Nevertheless, a search has been executed for these claims. The search has been based on the alleged effects of the compounds/compositions.

Box I.2

Present claims 1 and 22 relate to a compound defined by reference to a desirable characteristic or property, namely *having selective IK_{Ca} modulatory (inhibitory) activity*. The claims cover all compounds having this characteristic or property, whereas the application provides support within the meaning of Article 6 PCT and / or disclosure within the meaning of Article 5 PCT for only a very limited number of such compounds. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Independent of the above reasoning, the claims also lacks clarity (Article 6 PCT). An attempt is made to define the compound by reference to a result to be achieved. Further, the wording "which disease, disorder or condition relates to immune dysfunction" is not clear and concise. Again, this lack of clarity in the present case is such as to render a meaningful search over the whole of the claimed scope impossible. Consequently, the search has mainly been carried out for those parts of the claims which appear to be clear, supported and disclosed, namely those parts relating to the compounds in claim 18 and 1-[(2-chlorophenyl) diphenylmethyl]-1H-imidazole (clotrimazole) and closely related homologous compounds e.g mainly those compounds mentioned in the examples in the description at pages 22-24.

"The applicant's attention is drawn to the fact that claims relating to inventions in respect of which no international search report has been established will not be the subject of an international preliminary examination (Rule 66.1(e) PCT). This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure."

284498

INTERNATIONAL SEARCH REPORT
Information on patent family members

08/05/00

International application No.
PCT/DK 00/00253

Patent document cited in search report		Publication date		Patent family member(s)		Publication date
WO	9925347	A2	27/05/99	AU	1224599 A	07/06/99
WO	9734589	A1	25/09/97	AU BR CA EP US	2538897 A 9708230 A 2250092 A 0918514 A 6028103 A	10/10/97 04/01/00 25/09/97 02/06/99 22/02/00
WO	9418967	A1	01/09/94	AU CA EP JP US US US	6243494 A 2118313 A 0644760 A 8506594 T 5512591 A 5591763 A 5643936 A	14/09/94 01/09/94 29/03/95 16/07/96 30/04/96 07/01/97 01/07/97
US	5540931	A	30/07/96	WO CA EP US WO	9600058 A 2077647 A 0518872 A 4996193 A 9211860 A	04/01/96 08/07/92 23/12/92 26/02/91 23/07/92
WO	9601107	A1	18/01/96	AU AU WO	2880495 A 2880595 A 9601106 A	25/01/96 25/01/96 18/01/96